AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings of claims in the application:

LISTING OF CLAIMS:

Claims 1-8 (cancelled)

9. (new) A method for the treatment of HPV infections, comprising administering to a subject in need thereof an effective amount of human interferon in a liquid composition.

10. (new) The method according to claim 9, wherein the interferon is an amount of 100 to 500 IU/ml.

11. (new) The method according to claim 10, wherein the interferon is in an amount of 150 IU/ml.

12. (new) The method according to claim 9, wherein the human interferon is recombinant human interferon.

13. (new) The method according to claim 12, wherein the interferon is natural interferon.

14. (new) The method according to claim 9, wherein said liquid pharmaceutical composition is a water solution.

15. (new) A method for the treatment of infections of the general tract, comprising administering to a subject in need thereof an effective amount of human interferon in a liquid composition.

16. (new) A method for the treatment of warts or condylomatous lesions of the genital-tract mucosa, comprising administering to a subject in need thereof an effective amount of human interferon in a liquid composition.

17. (new) The method according to claim 9, wherein the composition is administered properly.

18. (new) The method according to claim 15, wherein the interferon is an amount of 100 to 500 IU/ml.

19. (new) The method according to claim 15, wherein the interferon is in an amount of 150 IU/ml.

20. (new) The method according to claim 15, wherein the human interferon is recombinant human interferon.

21. (new) The method according to claim 15, wherein the interferon is natural interferon.

22. (new) The method according to claim 15, wherein said liquid pharmaceutical composition is a water solution.

23. (new) The method according to claim 16, wherein the interferon is an amount of 100 to 500 IU/ml.

24. (new) The method according to claim 16, wherein the interferon is in an amount of 150 IU/ml.

25. (new) The method according to claim 16, wherein the human interferon is recombinant human interferon.

26. (new) The method according to claim 16, wherein the interferon is natural interferon.

27. (new) The method according to claim 16, wherein said liquid pharmaceutical composition is a water solution.

28. (new) The method according to claim 16, wherein the composition is administered properly.